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(54) Title: CULTURE MEDIA FOR IN VITRO FERTILIZATION AND EMBRYO TRANSFER

(57) Abstract

A culture medium for *in vitro* fertilization of human oocytes which comprises: NaCl (96.5 - 106.7 mM) KCl(4.46 - 4.92 mM) MgSO₄.7H₂O (0.18 - 0.22 mM) KH₂PO₄ (0.35 - 0.39 mM) CaCl₂.2H₂O (1.94 - 2.14 mM) NaHCO₃ (23.7 - 26.3 mM) Glucose (2.64 - 2.92 mM) Sodium Pyruvate (0.31 - 0.35 mM) Sodium Lactate (20.3 - 22.5 mM) Penicillin (95 - 105 units/ml) Phenol red (5 - 15 micrograms/ml). Culture media wherein the ratio of sodium ions to potassium ions is in the range of from 28 to 32 are preferred.

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CULTURE MEDIA FOR IN VITRO FERTILIZATION AND EMBRYO TRANSFER

The invention relates to culture media and more particularly to culture media for in vitro fertilization and embryo transfer.

The process of in vitro fertilization of human oocytes, cleavage of embryos and embryo transfer require that a culture medium be used to support the embryo for a period of up to three or four days during the various processes necessary for fertilization and early incubation before embryo transfer and reimplantation.

In the natural process of fertilization for a human oocyte the oocyte is supported within the mother within a fluid known as human tubal fluid and it is the object of the present invention to provide a culture medium as a synthetic human tubal fluid.

Approximation of the culture conditions as close as possible to those found in the natural environment of the gametes may be most likely to yield the best results. Using this rationale early workers have formulated a culture media similar in biochemical composition to human tubal fluid with varying rates of success. Examples of these include Tyrodes Medium T6, wm1 (Hoppe and Pitts), Modified Earles, and Hams F10.

One important characteristic of synthetic human tubal fluids appears to the ratio of sodium ions to potassium ions. For natural human tubal fluid this valve is approximately 18. Earlier attempts such as Tyrodes Medium T6 have a value of over 100. We have found that values in between these are most advantageous.

In the present invention we have devised a synthetic culture medium which is believed to approximate human tubal fluid but with desirable additional components and variations in the actual composition, including ratios of concentrations of sodium ions to potassium ions.

in one form therefore the present invention may be said to reside in a culture medium for in vitro fertilization and embryo transfer comprising the following compounds in the following ranges of concentration;

5	Sodium chloride (NaCl) Potassium chloride (KCl) Magnesium suiphate (MgSO47H2O)	96.5 - 106.7 mM 4.46 - 4.92mM 0.18 - 0.22mM
·	Potassium phosphate monobasic (KH ₂ PO ₄)	0.35 - 0.39mM
10	Calcium chloride 2 hydrate (CaCl ₂ 2H ₂ 0)	1.94 - 2.14mM
	Sodium bicarbonate (NaHCO3)	23:7 - 26.3mM
	Glucose	2.64 - 2.92mM
	Sodium Pyruvate	0.31 - 0.35mM
•	Sodium Lactate	20.3 - 22.5mM
15	Penicillin	95 - 105 units/m1
	Phenol red	5 - 15 micrograms/m1

In a preferred embodiment of the invention, the ratio of sodium ion concentration to potassium ion concentration is in the range from 28 to 32.

In a further preferred embodiment the ratio of concentrations of sodium ions to potassium ions is 29.3.

In one preferred embodiment of the invention the synthetic human tubal fluid may have the following composition;

25	Sodium Chloride (NaCl) Potassium chloride (KCl) Magnesium sulphate (MgSO ₄ 7H ₂ O)	101.6 mM 4.69 mM 0.20mM
25	Potassium phosphate monobasic (KH ₂ PO ₄)	0.37mM
·	Calcium chloride 2 hydrate (CaCl ₂ 2H ₂ 0)	2.04mM 25.0mM
30	Sodium bicarbonate (NaHCO3) Glucose	2.78mM

Sodium pyruvate		•	0.33mM
Sodium lactate	•	, 	21.4mM
Penicillin			100 units/ml
Phenol red			10 micrograms/mi

In another form the invention may be said to reside in a method of assisting with the in vitro fertilization of human oocytes including the step of handling the human oocytes in a culture medium, the culture medium being comprised of the compounds listed below in the range of compositions listed as follows;

10	Sodium chloride (NaC1)	96.5 - 106.7mM
	Potassium chloride (KC1)	4.46 - 4.92mM
	Magnesium sulphate (MgSO $_4$ 7H $_2$ O)	0.18 - 0.22mM
	Potassium phosphate monobasic (KH ₂ PO ₄)	0.35 - 0.39mM
15	Calcium chloride 2 hydrate	
•	(CaC1 ₂ 2H ₂ 0)	1.94 - 2.14mM
	Sodium bicarbonate (NaHCO3)	23.7 - 26.3mM
	Glucose	2.64 - 2.92mM
•	Sodium pyruvate	0.31 - 0.35mM
20	Sodium lactate	20.3 - 22.5mM
	Penicillin	95 - 105 units/ml
	Phenol red	5 - 15 micrograms/ml

In a preferred embodiment of this method of the invention, the ratio of sodium ion concentration to potassium ion concentration is in the range of from 28 to 32.

In a further preferred form of the invention, the ratio concentration of sodium ions to potassium ions is 29.3.

In a further preferred form of the invention, the method includes the step of handling the oocytes in a culture medium comprising;

Sodium Chloride (NaC1)

101.6 mM

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	Potassium chloride (Kcl)	4.69 mM
	Magnesium sulphate (MgSO $_4$ 7H $_2$ O)	0.20mM
	Potassium phosphate monobasic (KH ₂ PO ₄)	0.37mM
5	Calcium chloride 2 hydrate (CaCl ₂ 2H ₂ O)	2.04mM
	Sodium bicarbonate (NaHCO ₃)	25.0mM
•	Glucose	2.78mM
	Sodium pyruvate	0.33mM
10	Sodium lactate	21.4mM
	Penicillin	100 units/ml
	Phenol red	10 micrograms/ml

In a further form the invention may be said to reside in a culture medium for the in vitro fertilization of human oocytes including sodium potassium ions wherein the ratio of sodium ions to potassium ions is in the range of from 28 to 32.

in a preferred embodiment of this form of the invention, the ratio of sodium ions to potassium ions is 29.3.

This then generally describes the nature of the present invention and it will be seen that by this invention there is provided a culture medium which is not exactly the same as natural human tubal fluid, but which is capable of supporting in vitro fertilization.

To more clearly assist with the understanding of this invention reference will now be made to a preferred embodiment and tests to determine the efficacy of the preferred embodiment.

In one preferred embodiment synthetic human tubal fluid culture medium is as given in Table 1 below (marked synthetic HTF).

The medium may be prepared by using rainwater which has been distilled in glass six times. The bicarbonate-buffered medium is gassed for a minimum of five minutes with humidified 5% oxygen 5% carbon dioxide

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90% nitrogen mixture and sterilized by passage through a 0.45-0.2 micrometre filter membrane (Millipore, Sydney, Australia or Amicon Sterilet Adelaide, Australia) and then stored at 40°C for up to two weeks before use. A minimum of six hours or preferably the day before being used the bicarbonate buffered medium is gassed again for two to three minutes with the same gas mixture as above and the protein component is added.

In a similar way a known culture medium Tyrodes Medium T6 having a composition as given in Table 1 below was also prepared.

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10	COMPONENT	SYNTHETIC HFT	<u>16</u>
-	NaC1	101.6	99.4
	KC1	4.69	1.42
	MgSO ₄ .7H ₂ O	0.20	0.71
15	KH ₂ PO ₄	0.37	-
	CaC1 ₂ .2H ₂ 0	2.04	1.78
	NaHCO ₃	25	· 25 · ·
	Glucose	2.78	5.56
	Na pyruvate	0.33	0.47
20	Na lactate	21.4	24.9
	Penicillin	100 U/m1	100 U/m1
	Streptomycin SO ₄	-	50 ug/m1
	Phenol red	0.001% (w/v)	0.001% (w/v)

Tests have been carried out using both mouse embryo development in vitro and with initiation of human pregnancy in an endeavour to discover which components of the T6 medium and the synthetic human tubal fluid according to this invention might be responsible for observed differences in mouse embryo development in vitro and initiation of human pregnancies. The results show that for human pregnancy initiation almost three times as many pregnancies occurred when fertilization and culture were carried out

using the synthetic human tubal fluid of the present invention over the T6 medium.

In comparison of the two compositions a greatest difference in composition of the two media are their Na⁺/K⁺ ratios. We refer to these as sodium/potassium for the rest of the specification.

The ratios of concentrations of sodium ions to potassium ions for these are as follows:

Synthetic Human Tubal Fluid according to the present invention

29.3

· T6

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When media are tested with sodium/potassium levels varying from 150.5/1.42 millimoles to 148.2/5.06 millimoles results showed that there was significant linear and quadratic responses in the number of embryos developing to expanded blastocysts with increasing levels of K⁺.

in medium containing the potassium levels of T6 medium 75% of the zygotes developed which was significantly fewer than the 95-100% embryos developing when the potassium level was 2.3 to 5.1 millimoles which is the range for the synthetic human tubal fluid of the present invention.

The greatest number of mouse zygotes developing to expanded blastocysts when cultured in synthetic human tubal fluid medium of the present invention compared to T6 medium was paralleled by the three fold increase of the number of pregnancies initiated in those patients whose gametes had been fertilized and cultured in the medium in the present invention rather than T6 medium.

The present invention therefore provides a synthetic human tubal fluid culture medium which is more than just a direct replication of naturally occurring human tubal fluid but has enhanced viability.

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	Potassium chloride (Kcl) Magnesium sulphate (MgSO ₄ 7H ₂ O)	4.69 mM 0.20mM
	Potassium phosphate monobasic (KH ₂ PO ₄)	0.37mM
5	Calcium chloride 2 hydrate (CaCl ₂ 2H ₂ 0)	2.04mM
·	Sodium bicarbonate (NaHCO ₃)	25.0mM
	Glucose	2.78mM
•	Sodium pyruvate	0.33mM
10	Sodium lactate	21.4mM
	Penicillin	100 units/ml
	Phenol red	10 micrograms/ml

In a further form the invention may be said to reside in a culture medium for the in vitro fertilization of human oocytes including sodium potassium ions wherein the ratio of sodium ions to potassium ions is in the range of from 28 to 32.

In a preferred embodiment of this form of the invention, the ratio of sodium ions to potassium ions is 29.3.

This then generally describes the nature of the present invention and it will be seen that by this invention there is provided a culture medium which is not exactly the same as natural human tubal fluid, but which is capable of supporting in vitro fertilization.

To more clearly assist with the understanding of this invention reference will now be made to a preferred embodiment and tests to determine the efficacy of the preferred embodiment.

In one preferred embodiment synthetic human tubal fluid culture medium is as given in Table 1 below (marked synthetic HTF).

The medium may be prepared by using rainwater which has been distilled in glass six times. The bicarbonate-buffered medium is gassed for a minimum of five minutes with humidified 5% oxygen 5% carbon dioxide

90% nitrogen mixture and sterilized by passage through a 0.45-0.2 micrometre filter membrane (Millipore, Sydney, Australia or Amicon Sterilet Adelaide, Australia) and then stored at 40°C for up to two weeks before use. A minimum of six hours or preferably the day before being used the bicarbonate buffered medium is gassed again for two to three minutes with the same gas mixture as above and the protein component is added.

In a similar way a known culture medium Tyrodes Medium T6 having a composition as given in Table 1 below was also prepared.

TABLE 1

10	COMPONENT	SYNTHETIC HFT	<u>16</u>
	NaCl	101.6	99.4
	KC1	4.69	1.42
	MgS04.7H20	0.20	0.71
15	KH2P04	0.37	-
	CaC1 ₂ .2H ₂ O	2.04	1.78
	NaHCO ₃	25	25
	Glucose	2.78	5.56
	Na pyruvate	0.33	0.47
20	Na lactate	21.4	24.9
20	Penicillin	100 U/ml	100 U/m1
	Streptomycin SO ₄	-	50 ug/m1
	Phenol red	0.001% (w/v)	0.001% (w/v)

vitro and with initiation of human pregnancy in an endeavour to discover which components of the T6 medium and the synthetic human tubal fluid according to this invention might be responsible for observed differences in mouse embryo development in vitro and initiation of human pregnancies.

The results show that for human pregnancy initiation almost three times as many pregnancies occurred when fertilization and culture were carried out

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A culture medium for in vitro fertilization of human oocytes comprising:

5 .	Sodium chloride (NaCl) Potassium chloride (KCl) Magnesium sulphate (MgSO ₄ 7H ₂ O)	96.5 - 106.7 mM 4.46 - 4.92mM 0.18 - 0.22mM
· .	Potassium phosphate monobasic (KH ₂ PO ₄)	0.35 - 0.39mM
	Calcium chloride 2 hydrate (CaCl ₂ 2H ₂ O)	1.94 - 2.14mM
10	Sodium bicarbonate (NaHCO3)	23.7 - 26.3mM ·
	Glucose	2.64 - 2.92mM
	Sodium Pyruvate	0.31 - 0.35mM
	Sodium Lactate .	20.3 - 22.5mM
	Penicillin	95 - 105 units/m1
15	Phenol red	5 - 15 micrograms/mi

- 2. A culture medium in Claim 1 wherein the ratio of sodium ion concentration to potassium ion concentration is in the range of from 28 to 32.
- 3. A culture medium as in Claim 2 wherein the ratio of concentrations of sodium ions to potassium ions is approximately 29.3.
 - 4. A culture medium as in Claim 1 comprising approximately;

	·Sodium Chloride (NaCl)	101.6 mM
	Potassium chloride (KC1)	4.69 mM
	Magnesium sulphate (MgSO ₄ 7H ₂ O)	0.20mM
5	Potassium phosphate monobasic (KH ₂ PO ₄)	0.37mM
	Calcium chloride 2 hydrate	
	(CaCl ₂ 2H ₂ O)	2.04mM
	Sodium bicarbonate (NaHCO3)	25.0mM
LO	Glucose	2.78mM
	Sodium pyruvate	0.33mM

Sodium lactate	21.4mM	
Penicillin	100 units/ml	
Phenol red	10 micrograms/	ml

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5. A method of assisting with the in vitro fertilization of human oocytes including the steps of handling the oocytes in a culture medium, the culture medium being of the compounds being listed below in the range of compositions as follows;

5	Sodium chloride (NaCl)	96.5 - 106.7 mM
	Potassium chioride (KC1) Magnesium sulphate (MgSO47H2O)	4.46 - 4.92mM 0.18 - 0.22mM
	Potassium phosphate monobasic (KH ₂ PO ₄)	0.35 - 0.39mM
10	Calcium chloride 2 hydrate (CaC1 ₂ 2H ₂ O)	1.94 - 2.14mM
	Sodium bicarbonate (NaHCO3)	23.7 - 26.3mM
	Glucose	2.64 - 2.92mM
•	Sodium Pyruvate	0.31 - 0.35mM
15	Sodium Lactate	20.3 - 22.5mM
	Penicillin	95 - 105 units/m1
·	Phenol red	5 - 15 micrograms/m1

- 6. A method as in Claim 5 wherein the ratio of sodium ion concentration to potassium ion concentration is in the range of from 28 to 32.
- 7. A method as in Claim 6 wherein the ratio of concentration of sodium ions to potassium ions is approximately 29.3.
- 8. A method as in Claim 5 wherein the culture medium has a concentration as follows;

Sodium Chloride (NaCl)	101.6 mM
Potassium chloride (Kcl)	4.69 mM
Magnesium sulphate (MgSO ₄ 7H ₂ O)	0.20mM
Potassium phosphate	•
monobasic (KH ₂ PO ₄)	0.37mM

	Calcium chloride 2 hydrate (CaCl ₂ 2H ₂ O)	2.04mM
10	Sodium bicarbonate (NaHCO ₃)	25.0mM
	Glucose	2.78mM
	Sodium pyruvate	0.33mM
•	Sodium lactate	21.4mM
	Penicillin	100 units/mi
15	Phenol red	10 micrograms/ml

- 9. A culture medium for the in vitro fertilization of human oocytes including sodium and potassium ions wherein the ratio of sodium ions to potassium ions is in the range of from 28 to 32.
- 10. A culture medium as in Claim 9 wherein the ratio of sodium ions to potassium ions is approximately 29.3.

INTERNATIONAL SEARCH REPORT

INTERNATIONAL SEA	TRACH REPORT POT AU 86/00170 POT AU 86/00170
TO THE STATE OF SEVERAL CIRCULAR CONTRACTOR	symbols apply indicate alls 8
According to International Patent Classification (IPC) or to both National C	lassification and IPC
Int. C1.4 C12N 5/00, 5/02	
II FIELDS SEARCHED	
Minimum Documentation	fication Symbols
Classification System Classification System	
IPC C12N 5/00, 5/02	
Documentation Searched other than keep to the Extent that such Documents are in	Animum Documentation
AU : IPC as above	
III. DOCUMENTS CONSIDERED TO BE RELEVANT	Relevant passages 12 Relevant to Claim No 13
Category . Citation of Document, 11 with indication, where appropri	ate. of the following
P,X Fertility and Sterility, Volume October 1985, Quinn et al, "Imprin human in vitro fertilization medium based on the composition fluid', see pages 493-498	with the use of a
A 'In Vitro Fertilization and Emb Edited by A. Trounson and C. Woo (Churchill Livingstone) pages 3	2-33, 119-120
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1 agreeas (VEECH) 16 Janua	ary 1986 (16.01.86) 9-10
y US,A, 4473647 (CARPENTER et al. (25.09.84) See column 8 lines) 25 September 1984
X US.A. 4443432 (GARABEDIAN et a (17.04.84) See column 4 lines	1) 17 April 1984
 Special categories of cited documents: 19 "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cried to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "p" document published prior to the international filing date but leter than the priority date claimed 	"T" tater document published after the international filing dat or priority date and not in conflict with the application or cried to understand the principle or theory underlying in invention. "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered involve an inventive step. "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when it document is combined with one or more other such documents, such combination being obvious to a person said in the aft. "a" document member of the same patent family
IV. CERTIFICATION	Date of Mailing of this International Search Report
Date of the Actual Completion of the International Search 19 September 1986 (19.09.86)	29 SEP 1986
Australian Patent Office	G. MASTERS

ANNEX TO THE INTERNATIONAL SEARCH REPORT ON INTERNATIONAL APPLICATION NO. PCT/AU 86/00170

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report	Patent Family Members		
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US 4443432	AU 88532/82	CA 1187799	DK 4417/82
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END OF ANNEX